

(12) PATENT (11) Application No. AU 199891258 B2 (19) AUSTRALIAN PATENT OFFICE (10) Patent No. 735116 Title (54) Polynucleotide encoding a polypeptide having heparanase activity and expression of same in transduced cells  $(51)^7$ International Patent Classification(s) C12N 015/56 C12N 009/24 A61K 038/47 C12N 015/11 C12N 001/21 C12N 015/63 (21) Application No: 199891258 (22)Application Date: 1998.08.31 (87) WIPO No: WO99/11798 **Priority Data** (30)(31) (32) Date Number (33) Country 08/922170 1997.09.02 US 09/109386 1998.07.02 US (43) Publication Date: 1999.03.22 (43) Publication Journal Date: 1999.05.20 (44)Accepted Journal Date: 2001.06.28 (71) Applicant(s) Insight Strategy and Marketing Ltd.; Hadasit Medical Research Services and Development Ltd. (72)inventor(s) Iris Pecker; Israel Vlodavsky; Elena Feinstein (74) Agent/Attorney F.B. RICE and CO.,139 Rathdowne Street, CARLTON VIC 3053

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(54) Title: POLYNUCLEOTIDE ENCODING A POLYPEPTIDE HAVING HEPARANASE ACTIVITY AND EXPRESSION OF SAME IN TRANSDUCED CELLS

## (57) Abstract

A polynucleotide (hpa) encoding a polypeptide having heparanase activity, vectors including same, transduced cells expressing heparanase and a recombinant protein having heparanase activity.

The claims defining the invention are as follows:

- 1. A polynucleotide fragment comprising a polynucleotide sequence encoding a polypeptide having heparanase catalytic activity, wherein said polypeptide shares at least 70 % homology with SEQ ID NOs:10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
- 2. The polynucleotide fragment of claim 1, wherein said polynucleotide sequence includes nucleotides 63-1691 of SEQ ID NO:9, or nucleotides 139-1869 of SEQ ID NO:13.
- 3. The polynucleotide fragment of claim 1, wherein said polynucleotide sequence includes nucleotides 63-721 of SEQ ID NO:9.
- 4. The polynucleotide fragment of claim 1, wherein said polynucleotide is as set forth in SEQ ID NOs:9 or 13.
- 5. The polynucleotide fragment of claim 1, wherein said polynucleotide sequence includes a segment of SEQ ID NOs:9 or 13, said segment encodes said polypeptide having said heparanase catalytic activity.
- 6. The polynucleotide fragment of claim 1, wherein said polypeptide includes an amino acid sequence as set forth in SEQ ID NOs:10 or 14.
- 7. The polynucleotide fragment of claim 1, wherein said polypeptide includes a segment of SEQ ID NOs:10 or 14, said segment harbors said heparanase catalytic activity.
- 8. The polynucleotide fragment of claim 1, wherein said polynucleotide sequence is selected from the group consisting of double stranded DNA, single stranded DNA and RNA.
- 9. A polynucleotide fragment comprising a polynucleotide sequence at least 70 % homologous with SEQ ID NOs:9 or 13, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin, said polynucleotide sequence encoding a polypeptide having heparanase catalytic activity.

- 10). The polynucleotide fragment of claim 9, wherein said polynucleotide sequence is as set forth in SEQ ID NOs:9 or 13.
- 11. A vector comprising a polynucleotide sequence encoding a polypeptide having heparanase catalytic activity, wherein said polypeptide shares at least 70 % homology with SEQ ID NOs:10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
- 12. The vector of claim 11, wherein said polynucleotide sequence includes nucleotides 63-1691 of SEQ ID NO:9, or nucleotides 139-1869 of SEQ ID NO:13.
- 13. The vector of claim 11, wherein said polynucleotide sequence includes nucleotides 63-721 of SEQ 1D NO:9.
- 14. The vector of claim 11, wherein said polynucleotide sequence is as set forth in SEQ ID NOs:9 or 13.
- 15. The vector of claim 11, wherein said polynucleotide sequence includes a segment of SEQ ID NOs:9 or 13, said segment encodes said polypeptide having said heparanase catalytic activity.
- 16. The vector of claim 11, wherein said polypeptide includes an amino acid sequence as set forth in SEQ ID NOs:10 or 14.
- 17. The vector of claim 11, wherein said polypeptide includes a segment of SEQ ID NOs:10 or 14, said segment harbors said heparanase catalytic activity.
- 18. The vector of claim 11, wherein said polynucleotide sequence is selected from the group consisting of double stranded DNA, single stranded DNA and RNA.
  - 19. The vector of claim 11, wherein said vector is a baculovirus vector.
- 20. A vector comprising a polynucleotide sequence at least 70 % homologous with SEQ ID NOs:9 or 13, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin, said polynucleotide sequence encoding a polypeptide having heparanase catalytic activity.

- 21. The vector of claim 20, wherein said polynucleotide sequence is as set forth in SEQ ID NOs:9 or 13.
- 22. A host cell comprising an exogenous polynucleotide fragment including a polynucleotide sequence encoding a polypeptide having heparanase catalytic activity, wherein said polypeptide shares at least 70 % homology with SEQ ID NOs:10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
- 23. The host cell of claim 22, wherein said polynucleotide sequence includes nucleotides 63-1691 of SEQ ID NO:9, or nucleotides 139-1869 of SEQ ID NO:13.
- 24. The host cell of claim 22, wherein said polynucleotide sequence includes nucleotides 63-721 of SEQ 1D NO:9.
- 25. The host cell of claim 22, wherein said polynucleotide sequence is as set forth in SEQ ID NOs:9 or 13.
- 26. The host cell of claim 22, wherein said polynucleotide sequence includes a segment of SEQ ID NOs:9 or 13, said segment encodes said polypeptide having said heparanase catalytic activity.
- 27. The host cell of claim 22, wherein said polypeptide includes an amino acid sequence as set forth in SEQ ID NOs:10 or 14.
- 28. The host cell of claim 22, wherein said polypeptide includes a segment of SEQ ID NOs:10 or 14, said segment barbors said heparanase catalytic activity.
- 29. The host cell of claim 22, wherein said polynucleotide sequence is selected from the group consisting of double stranded DNA, single stranded DNA and RNA.
  - 30. The host cell of claim 22, wherein said cell is an insect cell.
- 31. A host cell comprising a polynucleotide sequence at least 70 % homologous with SEQ ID NOs:9 or 13, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group

(GCG) at the University of Wisconsin, said polynucleotide sequence encoding a polypeptide having heparanase catalytic activity.

- 32. The host cell of claim 31, wherein said polynucleotide sequence is as set forth in SEQ ID NOs:9 or 13.
- 33. A recombinant protein comprising a polypeptide having heparanase catalytic activity, said polypeptide shares at least 70 % homology with SEQ ID NOs:10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
- 34. The recombinant protein of claim 33, wherein said polypeptide includes a segment of SEQ ID NOs:10 or 14.
- 35. The recombinant protein of claim 33, wherein said polypeptide is as set forth in SEQ ID NOs:10 or 14.
  - 36. An amino acid sequence as set forth in SEQ ID NOs: 10 or 14.
  - 37. An amino acid sequence homologous to SEQ ID NOs:10 or 14.
- 38. A pharmaccutical composition comprising, as an active ingredient, a recombinant protein including a polypeptide having heparanase catalytic activity, said polypeptide shares at least 70 % homology with SEQ ID NOs:10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
- 39. The pharmaceutical composition of claim 38, wherein said polypeptide includes a segment of SEQ ID NOs:10 or 14.
- 40. The pharmaceutical composition of claim 38, wherein said polypeptide is as set forth in SEQ ID NOs: 10 or 14.
- 41. A modulator of heparin-binding growth factors, cellular responses to heparin-binding growth factors and cytokines, cell interaction with plasma lipoproteins, cellular susceptibility to viral, protozoa and bacterial infections or disintegration of neurodegenerative plaques comprising, as an active ingredient, a

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recombinant protein including a polypeptide having heparanase catalytic activity, said polypeptide shares at least 70% homology with SEQ ID NOs: 10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.

- 42. The modulator of claim 41, wherein said polypeptide includes a segment of SEQ ID NOs: 10 or 14 having heparanase catalytic activity.
- 10 43. The modulator of claim 41, wherein said polypeptide is as set forth in SEQ ID NOs: 10 or 14.
  - 44. A medical equipment comprising a medical device containing, as an active ingredient, a recombinant protein including a polypeptide having heparanase catalytic activity, said polypeptide shares at least 70% homology with SEQ ID NOs: 10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
  - 45. The medical equipment of claim 44, wherein said polypeptide includes a segment of SEQ ID NOs: 10 or 14.
    - 46. The medical equipment of claim 44, wherein said polypeptide is as set forth in SEQ ID NOs: 10 or 14.
    - 47. A host cell expressing a recombinant heparanase, wherein said recombinant heparanase shares at least 70% homology with SEQ ID NOs: 10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
    - 48. The host cell of claim 47, wherein said polypeptide includes a segment of SEQ ID NOs: 10 or 14.
    - 49. The host cell of claim 47, wherein said polypeptide is as set forth in SEQ ID NOs: 10 or 14.



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- 50. The host cell of claim 47, wherein said cell is an insect cell.
- 51. A cell extract or conditioned cell media or a partially purified cell extract or conditioned cell media comprising an extract or media of the host cell of any of claims 22-32 and 47-50 wherein said extract or media includes a polypeptide sharing at least 70% homology with SEQ ID NOs: 10 or 14.
- 52. A heparanase inhibitors screening system comprising the cell extract or extract or conditioned cell media or the partially purified cell extract or conditioned cell media of claim 51.
  - 53. A method of screening for a heparanase inhibitor, the method comprising assaying for heparanase catalytic activity of the recombinant protein of any of claims 33-35 in the presence and in the absence of, or in the presence of varying concentrations of, at least one compound being tested for a potential at inhibiting heparanase catalytic activity.
- overexpressing heparanase catalytic activity, wherein said heparanase catalytic activity is effected by a heparanase sharing at least 70% homology with SEQ ID NOs: 10 or 14, as determined using default parameters of a DNA sequence analysis software package developed by the Genetic Computer Group (GCG) at the University of Wisconsin.
  - 55. The system of claim 54, wherein said polypeptide includes a segment of SEQ ID NOs: 10 or 14 having heparanese catalytic activity.
- 56. The system of claim 54, wherein said polypeptide is as set forth in SEQ ID NOs: 10 or 14.
  - 57. A method of identifying a chromosome region harbouring a heparanase gene in a chromosome spread comprising the steps of:
- a) hybridizing the chromosome spread with a tagged polynucleotide
  35 probe at least 70% homologous with SEQ ID NOs: 9 or 13 as determined
  using default parameters of a DNA sequence analysis software package.



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developed by the Genetic Computer Group (GCG) at the University of Wisconsin;

- b) washing the chromosome spread, thereby removing excess of nonhybridized probe; and
- c) searching for signals associated with said hybridized tagged polynucleotide probe, wherein detected signals being indicative of a chromosome region harboring a heparanase gene.
- 58. A single stranded polynucleotide fragment comprising a polynucleotide sequence complementary to at least a portion of a polynucleotide strand defined by nucleotides 226 to 721 of SEQ ID NO: 9 having heparanese catalytic activity.
  - 59. A polynucleotide fragment according to any one of claims 1 to 10 substantially as hereinbefore described with particular reference to the examples.
  - 60. A vector according to any one of claims 11 to 21 substantially as hereinbefore described with particular reference to the examples.
  - 61. A host cell according to any one of claims 22 to 32 or 48 to 50 substantially as hereinbefore described with particular reference to the examples.
  - 62. A recombinant protein according to any one of claims 33 or 34 substantially as hereinbefore described with particular reference to the examples.
  - 63. A pharmaceutical composition according to any one of claims 38 or 40 substantially as hereinbefore described with particular reference to the examples.
    - 84. A modulator according to any one of claims 41 to 43 substantially as hereinbefore described with particular reference to the examples.



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- 65. Medical equipment according to any one of claims 44 to 46 substantially as hereinbefore described with particular reference to the examples.
- 5 66. A cell extract according to claim 51 substantially as hereinbefore described with particular reference to the examples.
  - 67. An inhibitor screen system according to claim 52 substantially as hereinbefore described with particular reference to the examples.
  - 68. An overexpression system according to any one of claims 54 to 56 substantially as hereinbefore described with particular reference to the examples.
- 15 69. A method according to claims 53 or 57 substantially as hereinbefore described with particular reference to the examples.

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